Docket No. UF-368XC1 Serial No. 10/809,631

## In the Claims

This listing of claims will replace all prior versions and listings of claims in this application.

1 (original). A method for treating idiopathic hyperhidrosis, wherein said method comprises administering to a patient a therapeutically effective amount of a 5-HT2C receptor activity affecting compound.

2 (original). The method of claim 1, wherein said 5-HT2C receptor activity affecting compound is selected from the group consisting of 5-HT2C receptor antagonists and 5-HT2C modulators.

3 (original). The method of claim 2, wherein said 5-HT2C receptor antagonist is selected from the group consisting of ketanserin, ritanserin, mianserin, meulergine, cyproheptadine, fluoxetine, mirtazapine, olanzapine, and ziprasidone.

4 (original). The method of claim 2, wherein said 5-HT2C modulator is selected from the group consisting of inverse agonists, partial agonists, and allosteric modulators.

5 (original). The method of claim 1, wherein said 5-HT2C receptor activity affecting compound is selected from the group consisting of (1R,2S,4R)-(-)-2-phenyl 2-(dimethylaminoethoxy)-1,7,7-trimethyl-bicyclo[2.2.1]heptane and (1R,2S,4R)-(-)-2-phenyl-2-(methylaminoethoxy)-1,7,7-trimethyl-bicyclo[2.2.1]heptane.

6 (original). The method of claim 1, wherein said 5-HT2C receptor activity affecting compound is administered to the patient via a route selected from the group consisting of oral, topical, mucosal, systemic, parenteral, intravenous, intraperitoneal, subcutaneous, intramuscular, intraoral, rectal, epicutaneous, transdermal, intranasal, sublingual, buccal, intradural, intraocular, intrarespiratory, and intra nasal inhalation.

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7 (original). The method of claim 1, wherein said 5-HT2C receptor activity affecting compound is administered to the patient via liposome delivery systems.

8 (original). The method of claim 1, further comprising the step of concurrently administering an agent used to treat sweating.

9 (original). The method of claim 8, wherein said agent is selected from the group consisting of antiperspirants, acetylcholine-blocking compounds, and beta blockers.

10 (original). The method of claim 9, wherein said agent is selected from the group consisting of aluminum acetate, aluminum sulfate, aluminum chloride, propranolol, glycopyrrolate, atropine, propantheline bromide, and oxybutynin.

11 (original). The method of claim 1, further comprising the step of concurrently administering a method for treating sweating.

12 (original). The method of claim 11, wherein said method for treating sweating is selected from the group consisting of iontophoresis, endoscopic thoracic sympathicotomy, and botulinum toxin injection.

13 (original). A method for treating symptoms or associated conditions of idiopathic hyperhidrosis, wherein said method comprises administering to a patient a therapeutically effective amount of a 5-HT2C receptor activity affecting compound.

14 (original). The method of claim 13, wherein said 5-HT2C receptor activity affecting compound is selected from the group consisting of 5-HT2C receptor antagonists and 5-HT2C modulators.

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15 (original). The method of claim 14, wherein said 5-HT2C receptor antagonist is selected from the group consisting of ketanserin, ritanserin, mianserin, meulergine, cyproheptadine, fluoxetine, mirtazapine, olanzapine, and ziprasidone.

16 (original). The method of claim 14, wherein said 5-HT2C modulator is selected from the group consisting of inverse agonists, partial agonists, and allosteric modulators.

17 (original). The method of claim 13, wherein said 5-HT2C receptor activity affecting compound is selected from the group consisting of (1R,2S,4R)-(-)-2-phenyl 2-(dimethylaminoethoxy)-1,7,7-trimethyl-bicyclo[2.2.1]heptane and (1R,2S,4R)-(-)-2-phenyl-2-(methylaminoethoxy)-1,7,7-trimethyl-bicyclo[2.2.1]heptane.

18 (original). The method of claim 13, wherein said 5-HT2C receptor activity affecting compound is administered to the patient via a route selected from the group consisting of oral, topical, mucosal, systemic, parenteral, intravenous, intraperitoneal, subcutaneous, intramuscular, intraoral, rectal, epicutaneous, transdermal, intranasal, sublingual, buccal, intradural, intraocular, intrarespiratory, and intra nasal inhalation forms.

19 (original). The method of claim 13, wherein said 5-HT2C receptor activity affecting compound is administered to the patient via liposome delivery systems.

20 (original). The method of claim 13, further comprising the step of concurrently administering an agent used to treat sweating.

21 (original). The method of claim 20, wherein said agent is selected from the group consisting of antiperspirants, acetylcholine-blocking compounds, and beta blockers.

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22 (original). The method of claim 21, wherein said agent is selected from the group consisting of aluminum acetate, aluminum sulfate, aluminum chloride, propranolol, glycopyrrolate, atropine, propantheline bromide, and oxybutynin.

23 (original). The method of claim 13, further comprising the step of concurrently administering a method for treating sweating.

24 (original). The method of claim 23, wherein said method for treating sweating is selected from the group consisting of iontophoresis, endoscopic thoracic sympathicotomy, and botulinum toxin injection.

25-31 (canceled).

32 (original). A method for prophylactically preventing or minimizing perspiring, wherein said method comprises administering to a patient a therapeutically effective amount of a 5-HT2C receptor activity affecting compound.

33 (original). The method of claim 32, wherein said 5-HT2C receptor activity affecting compound is selected from the group consisting of 5-HT2C receptor antagonists and 5-HT2C modulators.

34 (original). The method of claim 33, wherein said 5-HT2C receptor antagonist is selected from the group consisting of ketanserin, ritanserin, mianserin, meulergine, cyproheptadine, fluoxetine, mirtazapine, olanzapine, and ziprasidone.

35 (original). The method of claim 33, wherein said 5-HT2C modulator is selected from the group consisting of inverse agonists, partial agonists, and allosteric modulators.

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36 (original). The method of claim 32, wherein said 5-HT2C receptor activity affecting compound is selected from the group consisting of (1R,2S,4R)-(-)-2-phenyl 2-(dimethylaminoethoxy)-1,7,7-trimethyl-bicyclo[2.2.1]heptane and (1R,2S,4R)-(-)-2-phenyl-2-(methylaminoethoxy)-1,7,7-trimethyl-bicyclo[2.2.1]heptane.

37 (original). The method of claim 32, wherein said 5-HT2C receptor activity affecting compound is administered to the patient via a route selected from the group consisting of oral, topical, mucosal, systemic, parenteral, intravenous, intraperitoneal, subcutaneous, intramuscular, intraoral, rectal, epicutaneous, transdermal, intranasal, sublingual, buccal, intradural, intraocular, intrarespiratory, and intra nasal inhalation forms.

38 (original). The method of claim 32, wherein said 5-HT2C receptor activity affecting compound is administered to the patient via liposome delivery systems.

- 39 (original). The method of claim 32, further comprising the step of concurrently administering an agent used to treat sweating.
- 40 (original). The method of claim 39, wherein said agent is selected from the group consisting of antiperspirants, acetylcholine-blocking compounds, and beta blockers.
- 41 (original). The method of claim 40, wherein said agent is selected from the group consisting of aluminum acetate, aluminum sulfate, aluminum chloride, propranolol, glycopyrrolate, atropine, propantheline bromide, and oxybutynin.
- 42 (original). The method of claim 32, further comprising the step of concurrently administering a method for treating sweating.

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43 (original). The method of claim 32, wherein said method for treating sweating is selected from the group consisting of iontophoresis, endoscopic thoracic sympathicotomy, and botulinum toxin injection.

44 (new). The method of claim 13, wherein the associated conditions of idiopathic hyperhidrosis are selected from the group consisting of: eczematous dermatitis; contact dermatitis; miliaria; and bromhidroses.

45 (new). The method of claim 6, wherein said receptor activity affecting compound is administered to the patient via injection.

46 (new). The method of claim 18, wherein said receptor activity affecting compound is administered to the patient via injection.

47 (new). The method of claim 37, wherein said receptor activity affecting compound is administered to the patient via injection.